

667070" 20488260

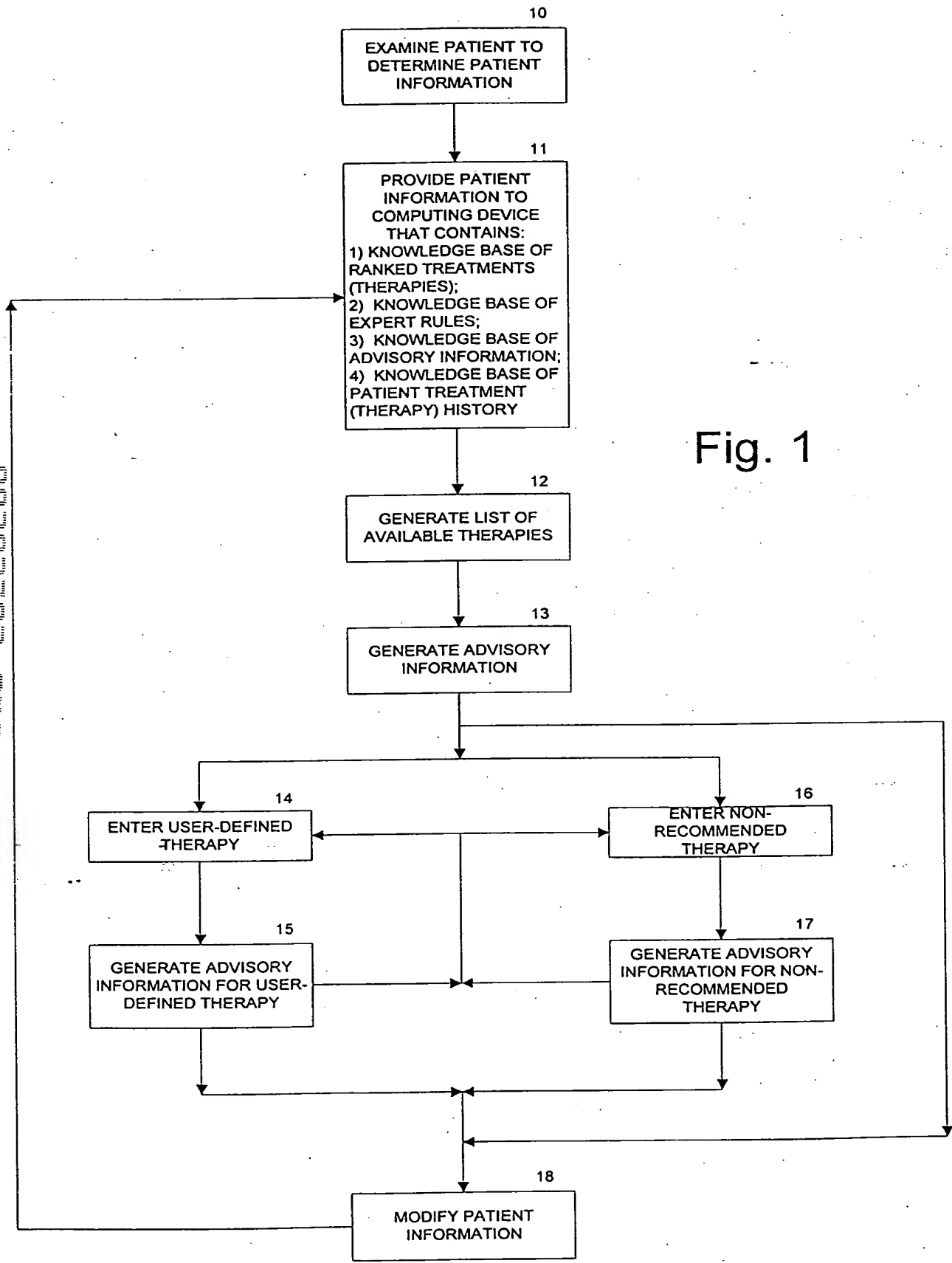


Fig. 1

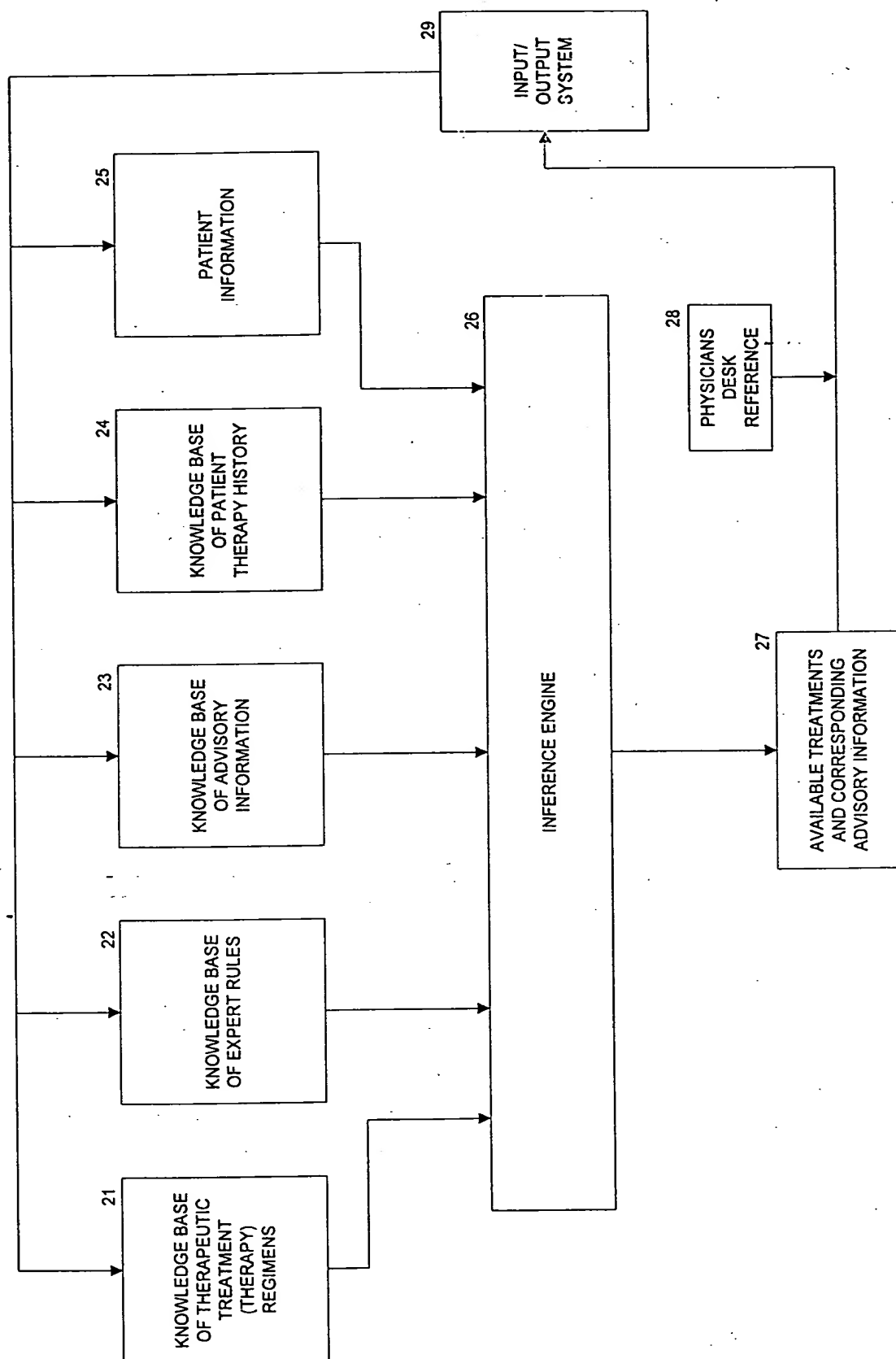


Fig. 2

667040" 20263260

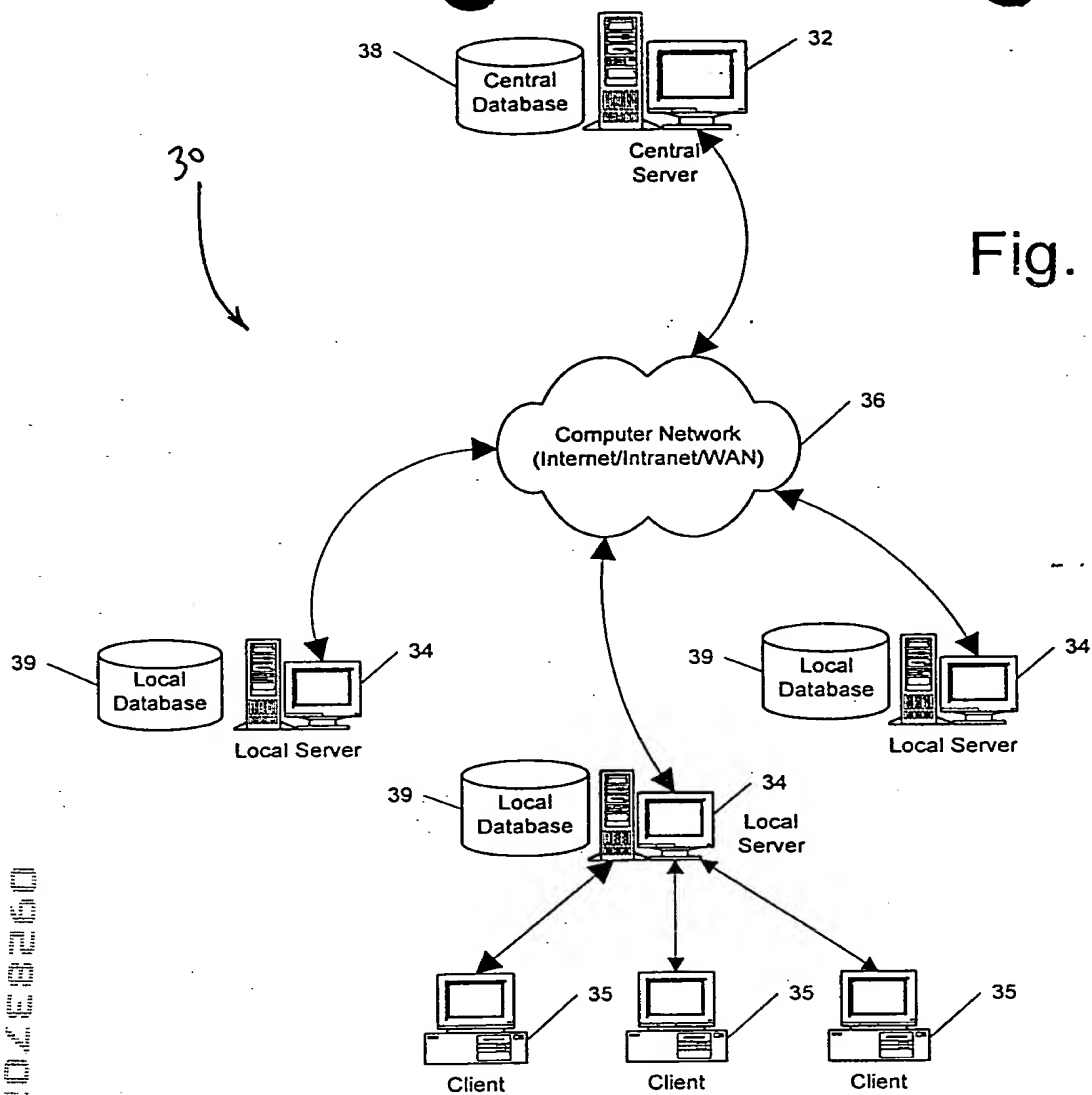


Fig. 3

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Medical History		Chart		Therapy Evaluation	
General		Pin		Save	
Patient Id		IPMS Number		Weight (kg)	
Birth Date		Copy		Can Take Solid Dosage	
And Viral Load		Specimen Date		Specimen Date	
Viral Load (copies/ml)		Value		Prev Value	
HIV-1 RNA		Specimen Date		Value(s)	
Phenotypic		Specimen Date		Value(s)	
ABV-1 RNA		Specimen Date		Value(s)	
Intolera		Specimen Date		Value(s)	
Hemoglobin		Specimen Date		Value (g/dL)	
Neutrophils		Specimen Date		Value (cells/cubic mm)	
Hepatic Function		Specimen Date		Value (ALT/SGPT (U/L))	
Renal Function		Specimen Date		Value (Sodium Creatinine)	

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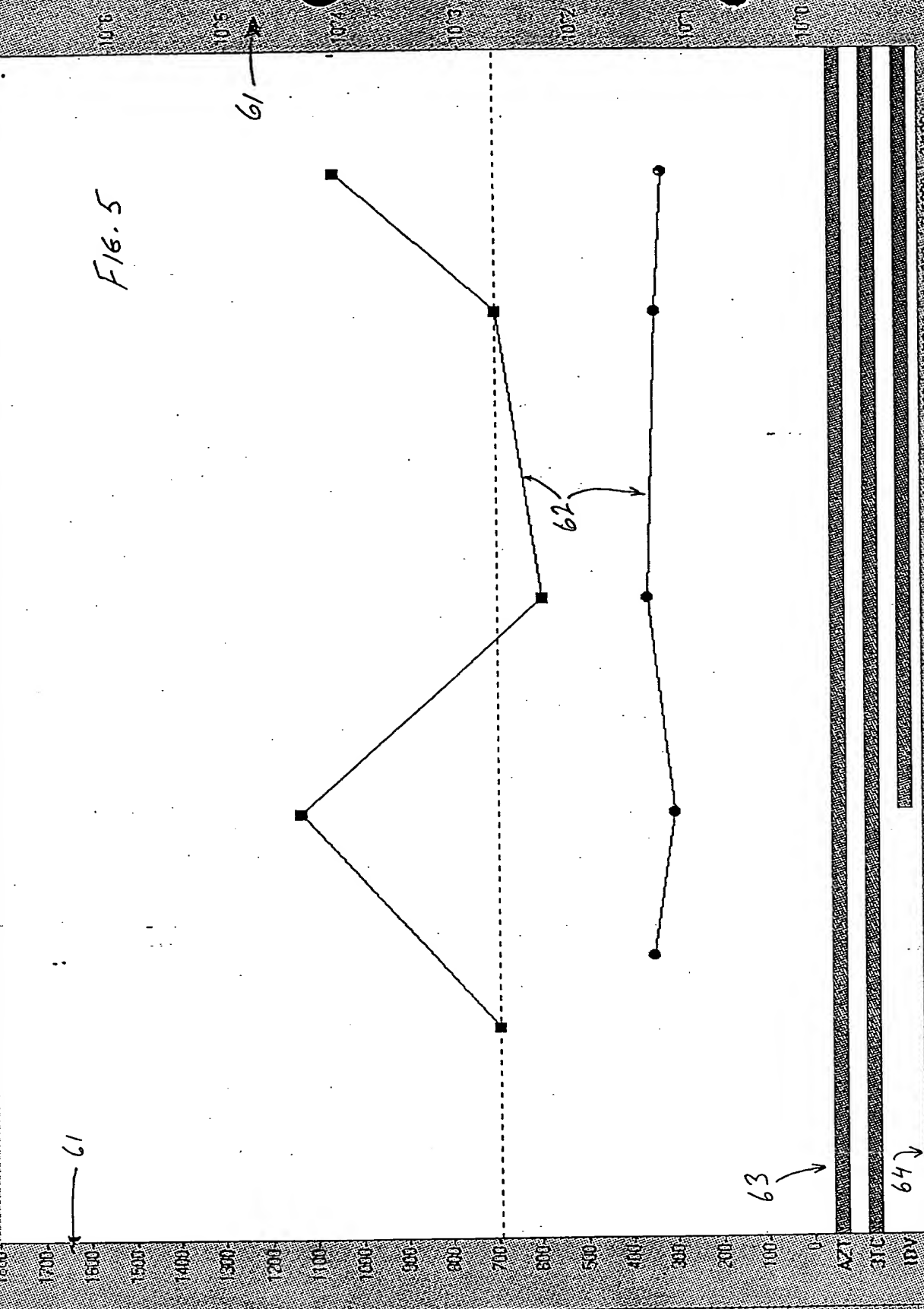
FIG. 4

60a 70a

TPMS Patient

Medical History Chart Therapy Evaluation

CD4 (cells/mm<sup>3</sup>) Viral Load (copies/ml)



12/1997 1/1998 2/1998 3/1998 4/1998 5/1998 6/1998 7/1998 8/1998 9/1998 10/1998 11/1998 12/1998 1/1999 2/1999 3/1999








AZT 3TC IDV

TPMS



092810 104199  
66T040 2078260

F16.7

Icon	Meaning
	Indicates that there were no critical alerts for the therapy, however, general warnings and advisories should be read in the Therapy Details box.
	Indicates that there were no critical alerts for the therapy, however, general warnings and advisories should be read in the Therapy Details box. The book indicates that therapy has been studied and a reference is available to review.
	Indicates a yellow alert. There is important information about this therapy that must be reviewed.
	Indicates a yellow alert. There is important information about this therapy that must be reviewed. The book indicates that therapy has been studied and a reference is available to review.
	Indicates a red alert, which means critical and possible life-threatening situation may exist or may be created with this therapy. Information in the Therapy Details section must be read for this therapy to be considered.
	Indicates a red alert, which means critical and possible life-threatening situation may exist or may be created with this therapy. Information in the Therapy Details section must be read for this therapy to be considered. The book indicates that therapy has been studied and a reference is available to review.
	Indicates the therapy is not recommended.

F16-8

73a 78 73b

**Alert Patient** | **Medical History** | **Chemotherapy Evaluation** | **Life at Current (New)** | **Stop Therapy**

**Therapy Being Evaluated:** AZT, ddI, 3TC, RITV

**73c** **STOP DRUG INTERACTION RED ALERT STOP**

*Read the following Red Drug Contra-Indication Alerts for this therapy:*

**Drug Interactions Alert:** Patient is currently taking cisplatin, co-administration of Norvir (Ritonavir/RTV) with certain non-sedating antihistamines, sedative hypnotics, or antiarrhythmics may result in potentially serious and/or life-threatening adverse events due to possible effects of Norvir (Ritonavir/RTV) on the hepatic metabolism of certain drugs. Norvir (Ritonavir/RTV) can produce large increases in plasma concentrations of certain highly metabolized drugs. Norvir (Ritonavir/RTV) should not be coadministered with upiroxalen, amiodarone, astemizole, bepridil, bupropion, cispapine, clozapine, clobazepam, encainide, flucanazole, flutazepam, meprobamate, midazolam, propafenone, propoxyphene, quinidine, rifabutin, rifampin, terfenadine, triazolam or zolpidem. Patient is taking cispapine and in order to use this therapy, that drug should be replaced with a non-contraindicated substitute. *See DIL, Commentary 23*

**73d** **Dosages**

- Ritonavir 300mg q12h (2 pills/day, \$9.56/day)
- Videx 125mg q12h (4 pills/day, \$4.22/day)
- Inivase 400mg q12h, taken within 2 hours after a full meal (4 pills/day, \$3.47/day)
- Norvir 400mg q12h (2 pills/day, \$1.43/day)

(# indicates adjusted dosage)

**73e** **Dosage Adjustments:** The following dosage adjustments messages apply to this therapy:

- Dosage Notice: This therapy contains both zalcitabine and zalcitabine. When zalcitabine and zalcitabine are used together the dosage of each drug is reduced by 1/3. The dosage for these drugs has been set accordingly. *See DIL, Commentary 23*

**73f** **Inivase (zalcitabine/SQV):** The following Warnings and Advisories apply to Inivase (zalcitabine/SQV):

- Drug Interactions Information: Compounds that are substrates of CYP3A4 (e.g., calcium channel blockers, clindamycin, dapsone, quinidine, triazolam) may have elevated plasma concentrations when coadministered with Inivase (zalcitabine/SQV); therefore, patients should be monitored for toxicities associated with such drugs when taking Inivase (zalcitabine/SQV). *See DIL, Commentary 23*

73

73d

73e

73f

73g



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Therapy Options			
Therapy	Eff	Adap	Safety
1 d4T, 3TC, IDV	1	1	
1 AZT, 3TC, IDV	1	1	
1 d4T, 3TC, NFV	1	1	
1 AZT, 3TC, NFV			
1 d4T, 3TC, NFV			
1 AZT, 3TC, NFV			
1 ddI, d4T, 3TC			
1 d4T, 3TC, NFV			
1 d4T, 3TC, NFV			

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FIG. 9

Therapy-B  
Evaluated

General

- Vi
- Me

Show Abstract for Retrovir

Show Abstract for Epivir

Show Abstract for Viracept

Show Therapy Study

Print Details for AZT, 3TC, NFV

Print Top 10 Therapy Option Details

Hide Column "Eff"

Hide Column "Adap"

Hide Column "Safety Considerations"

Show Column "Med"

Show Column "Drug"

Hide Column "Freq"

Hide Column "Pills"

Hide Column "Cost"

General		Patient Information		Comments/Prognosis	
Field	Value	Field	Value	Field	Value
Patient ID	demo1	Birth Date	1/1/1960	TPMS Number	
Physician		Gender	Male	Weight (kg)	55.00
CD4 and Viral Load		AIDS Diagnosis		AIDS Defining Event	
CD4 (cells/cubic mm)	320	Date		Current ARV Therapy	
Current Viral Load	12000	Specimen Date	3/1/1999	Start Date	
Previous Viral Load	500	VI Units	C/mL	End Date	
HIV Genotype		Non-ARV Drugs		Therapy Drug	
Phenotype		AZT, 3TC, IDV		tobramycin	
Allergy/Hyper		Therapy Drug		Start Date	
Intolerance		tobramycin		1/1/1999	
Hemoglobin		Neutropathy		Value	
Specimen Date	3/1/1999	Value (g/dL)	12.00	Date	3/1/1999
Neutrophils		Pancreatitis		Value	
Specimen Date	3/1/1999	Cells/cubic mm	1500	Date	3/1/1999
Hepatic Function		Renal Function		Value	
Specimen Date	3/1/1999	AST/SGOT (U/L)	49	Specimen Date	3/1/1999
		ALT/SGPT (U/L)	45		
Serum Creatinine		Diagnosis		Est. Creatinine	
3/1/1999	2.00	No			39.39

FIG-10A

54b

F1

54a

F2



660

60a 70a

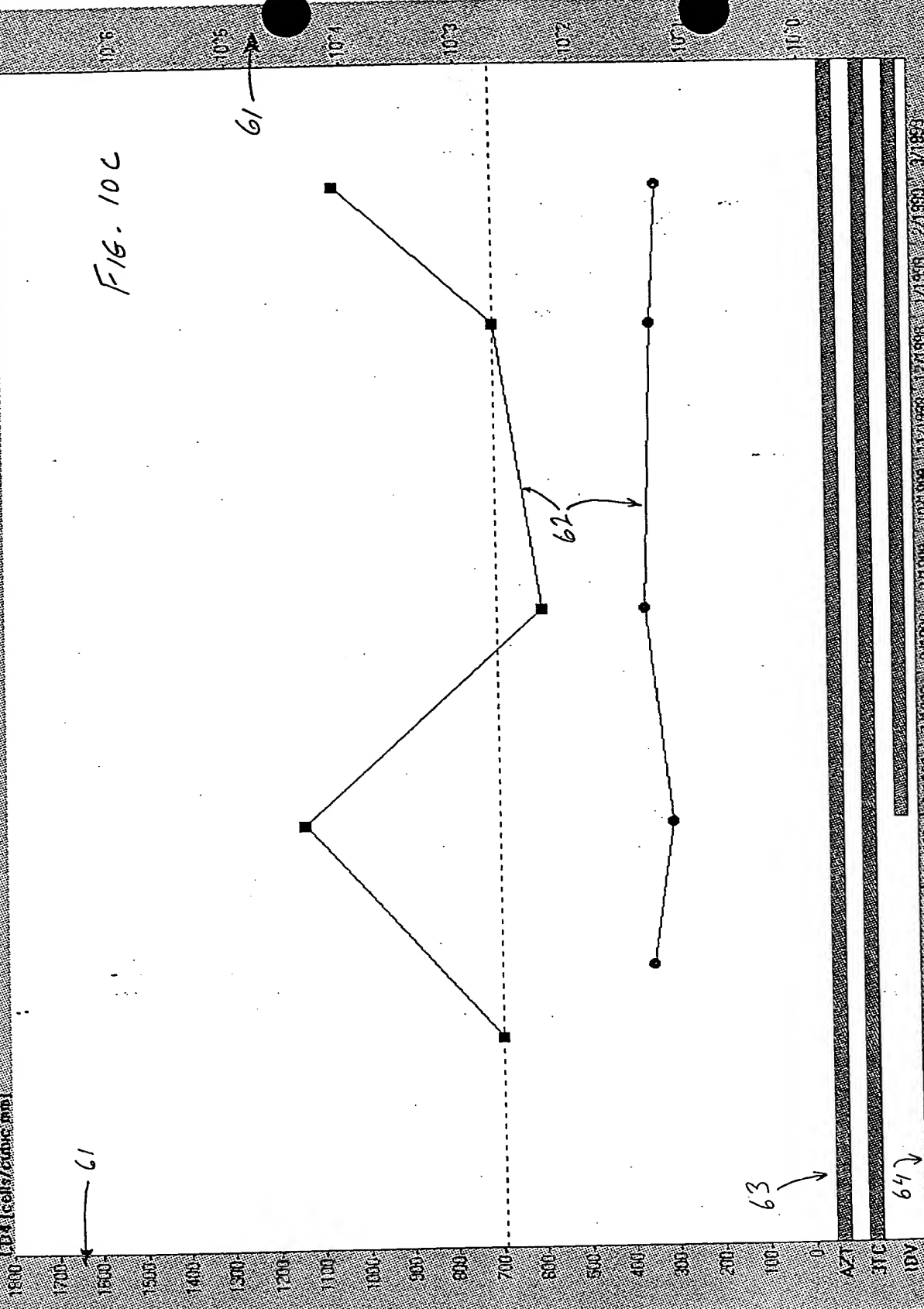
TPMS Patient

Medical History Chan Therapy Evaluation

Viral Load (copies/mL)

CD4 (cells/cubic mm)

FIG. 10C



63

64

2/1997 1/1996 2/1996 3/1996 4/1996 5/1996 6/1996 7/1996 8/1996 9/1996 10/1996 11/1996 12/1996 1/1997 2/1997 3/1997 4/1997 5/1997 6/1997 7/1997 8/1997 9/1997 10/1997 11/1997 12/1997 1/1998 2/1998 3/1998 4/1998 5/1998 6/1998 7/1998 8/1998 9/1998 10/1998 11/1998 12/1998 1/1999 2/1999 3/1999

TPMS +1

Navigation icons: Home, Back, Forward, Stop, Print, Web, Mail, etc.



70

Fig. 10D

M81

TPMS Patient

Medical History

Chart

Therapy Evaluation

General

Patient Id

demo1

Birth Date

1/1/1960

Gender

Male

TPMS Number

Weight (kg)

55.00

Solid Doseage

Yes

Empty

Current Popus

Save

Print

EDA and Viral Load

EDA

3/3/1999

Current Viral Load

55.00

Previous Viral Load

55.00

HIV Genotype

Specimen Date

3/3/1999

Phenotype

Specimen Date

3/3/1999

Allergy/Hyper

Specimen Date

3/3/1999

Intolerance

Specimen Date

3/3/1999

Hemoglobin

Specimen Date

3/3/1999

Neutrophils

Specimen Date

3/3/1999

Hepatic Function

Specimen Date

3/3/1999

Boundry and Prequalification Messages

Please be aware that the following boundry and prequalification conditions currently apply to the patient.

• Poor Viral Suppression  $\Delta$ : The patient's viral load count either did not decrease  $\geq 5$  log from the last point or is not below the viral load reduction goal. Unless lab error is at fault, consider changing therapy. More Info PQ! P=QualA6, Commentary445

Data Needed Soon - Caution

• No Baseline Viral Load Value: Please specify which viral load value or values (an average of two points) you wish to be set as the baseline viral load value for this patient.

BoundryZY, Commentary41a

TPMS +1

150

60a 70a

FIG. 11A

TPMS Patient		Medical History		Chart		Therapy Evaluation	
General		Patient ID: ARV naive		Birth Date: 1/5/1968		TPMS Number: [ ]	
Physician: [ ]		Gender: Male		First: [ ]		Save	
CD4 and Viral Load		Specimen Date		Value		Specimen Date	
CD4 (cells/cubic mm)		2/20/1999		350		2/20/1999	
Current Viral Load		2/20/1999		31000		C/mL	
Previous Viral Load		12/29/1998		19000		C/mL	
HIV Genotype		Specimen Date		Value		Specimen Date	
Phenotype		2/1/1999		[ ]		[ ]	
Allergy/Hyper		2/1/1999		[ ]		[ ]	
Intolerance		2/1/1999		[ ]		[ ]	
Hemoglobin		Specimen Date		Value (g/dL)		Specimen Date	
Neutrophils		2/1/1999		12.50		[ ]	
Neutrophils		Specimen Date		Value (cells/mm <sup>3</sup> )		Specimen Date	
Neutrophils		2/1/1999		1350		[ ]	
Hepatic Function		Specimen Date		AST/SGOT (U/L)		Specimen Date	
Hepatic Function		2/1/1999		35		[ ]	
Renal Function		Specimen Date		ALT/SGPT (U/L)		Specimen Date	
Renal Function		2/1/1999		35		[ ]	
AIDS Diagnostic		Date		AIDS Diagnostic Event		Date	
AIDS Diagnostic		2/1/1999		[ ]		[ ]	
Current ARV Therapy		V		D		X	
Non-ARV Drugs		Therapy Drug		Route		Start Date	
Non-ARV Drugs		Prozac Pulvules & Liquid, O...		oral		10/5/1998	
Non-ARV Drugs		Bacilum DS Tablets		oral		12/8/1998	
Weight (kg)		Date		Value		Date	
Weight (kg)		2/1/1999		73.00		[ ]	
Solid Dosage		Date		Value		Date	
Solid Dosage		2/1/1999		Yes		[ ]	



70a

TPMS Patient

Medical History Chart Therapy Evaluation

General  
Patient ID: ARV naive  
Birth Date: 1/5/1968  
Gender: Male  
TPMS Number:   
Weight (kg):   
Solid Doseage:   
Date: 2/1/1999  
Value: 73.00  
Yes  
OK  
Cancel

CD4 and Viral Load  
CD4 (cells/mm<sup>3</sup>): 27  
Current Viral Load: 27  
Previous Viral Load: 12  
HIV Genotype:   
Phenotype:   
Allergy/Hyper:   
Intolerance:   
Hemoglobin:   
Neutrophils:   
Hepatic Function:   
Specimen Date: 2/1/1999  
Value: 35

Boundry and Prequalification Messages  
Please be aware that the following boundry and prequalification conditions currently apply to this patient.  
Therapy Initiation: Current treatment guidelines recommend initiation of antiRetroviral therapy for HIV -infected patients with HIV RNA (viral load) concentrations greater than 20,000 copies/ml (10,000 E<sub>q</sub>/ml bDNA) or CD4 counts less than 300 cells/uL (Ann. Int. Med., 1998). PreQualM, Commentary61  
Combination Therapy Recommended: Experts agree that the goal of antiRetroviral therapy should be to reduce the viral load to as low a level as possible for as long as possible. Initiation of therapy with a combination containing 2 nucleoside reverse transcriptase inhibitors (NRTI's) and a potent protease inhibitor have been shown to provide enhanced clinical benefit versus 2 drug combinations with regard to reduction in viral load and improved clinical outcomes. PreQualM, Commentary66

m32

Fig-113

Therapy	Eff	Ad	Safety Considerations	Freq	Cost
<input checked="" type="radio"/> AZT, ddI, 3TC, SQV-SGC	1	1		q8h	\$43.46
<input checked="" type="radio"/> ddI, 3TC, NFV	1	1		q8h	\$34.78
<input checked="" type="radio"/> AZT, 3TC, IDV	1	1		q8h	\$32.24
<input checked="" type="radio"/> AZT, 3TC, NFV	1	1		q8h	\$35.81
<input checked="" type="radio"/> ddI, 3TC, IDV	1	1		q8h	\$31.20
<input checked="" type="radio"/> AZT, ddI, RTV, DLV	2	2	DLV+RTV	q8h	\$45.99
<input checked="" type="radio"/> ddI, ddI, IDV, NVP	2	2		q8h	\$42.55
<input checked="" type="radio"/> ddI, 3TC, RTV	2	2		q12h	\$38.46
<input checked="" type="radio"/> AZT, ddI, RTV, NVP	2	2		q12h	\$47.10

See More See All Top 10 Full Screen Evaluation

Therapy Being Evaluated None

Antiretroviral Drug

Nucleoside Analogues (NRTI)

☐ AZT (Retrovir/zidovudine)

☐ ddI (Videx/ddanosine)

☐ ddC (Hivid/zalcitabine)

☐ 3TC (Epivir/lamivudine)

☐ d4T (Zenit/stavudine)

☐ ABC (Ziagen/abacavir)

Protease Inhibitors (PI)

☐ IDV (Crivari/indinavir)

☐ SQV-HGC (Invirase/sequinavir)

☐ SQV-SGC (Entovase/sarinavir)

Usage of Current Therapy

- **WARNING:** Before initiating any antiRetroviral treatment regimen, the complete product information for each therapeutic component should be consulted.
- **Viral Load Testing Required:** Viral load testing should be repeated 21-35 days after initiation of, or a change of, antiRetroviral therapy to evaluate therapeutic efficacy and patient compliance. CmtGenY, Commentary65
- **Therapy Initiation:** Current treatment guidelines recommend initiation of antiRetroviral therapy for HIV-infected patients with HIV RNA (viral load) concentrations greater than 20,000 copies/ml (10,000 E<sub>q</sub>/ml bDNA) or CD4 counts less than 500 cells/ $\mu$ L (Ann. Int. Med., 1998). PreQualM, Commentary61
- **Combination Therapy Recommended:** Experts agree that the goal of antiRetroviral therapy should be to reduce the viral load to as low a level as possible for as long as possible. Initiation of therapy with a combination containing 2 nucleoside reverse transcriptase inhibitors (NRTIs) and a potent protease inhibitor have been shown to provide enhanced clinical benefit versus 2 drug combinations with regard to reduction in viral load and improved clinical outcomes. PreQualM, Commentary66



Therapy Being Evaluated  
AZT ddi RTV DLV

## Recommended Dosages

- Retrovir 300mg q12h (2 pills/day, \$9.56/day)
- Videx 200mg q12h (4 pills/day, \$6.78/day)
- Norvir 600mg q12h (12 pills/day, \$22.26/day)
- Rescriptor 400mg q8h (12 pills/day, \$7.39/day)

- AZT: Interrupt Retrovir if anemia and/or neutropenia develops. More Info 036 DosGenA, Commentary36

- ddi: When treatment with other drugs known to cause pancreatic toxicity is required (for example, IV pentamidine), suspension of Videx should be considered.

CmtGenA, Commentary13

- ddi: If patients develop symptoms of neuropathy, Videx therapy should be interrupted. DosGenB, Commentary40

- ddi: Clinical signs suggestive of pancreatitis should prompt dose suspension of Videx and careful evaluation of the possibility of pancreatitis. Only after pancreatitis has been ruled out should dosing be resumed. DosGenB, Commentary39

- DLV: Skin rash attributable to Rescriptor may occur during first 21 days. More Info 054 CmtGenS, Commentary54

- ddi: Videx should not be administered with a prescription antibiotic containing any form of tetracycline. CmtGenA, Commentary15

- ddi: Plasma concentrations of some quinolone antibiotics are decreased when administered with antacids containing magnesium or aluminum. Therefore, doses of quinolone antibiotics should not be administered within 2 hours of taking Videx. CmtGenA, Commentary16

- RTV: Monitor for decreased AUC of Norvir and associated adverse events when concomitant with use of drugs that increase CYP3A activity (including tobacco). More Info 026 CmtGenH, Commentary26

TPMS Patient

Medical History

Chart

Therapy Evaluation

Evaluate Current Therapy

None

Show 1 Drug Therapies

Show 2 Drug Therapies

Show 3 Drug Therapies

Show 4 Drug Therapies

Show 5 Drug Therapies

Antiretroviral Drugs

Nucleoside Analogues (NRTI)

☐ AZT (Retrovir/zidovudine)
 ☐ ddI (Videx/ddanosine)
 ☐ ddC (Hivid/zalcitabine)
 ☐ 3TC (Epivir/lamivudine)
 ☐ d4T (Zenit/stavudine)
 ☐ ABC (Ziagen/abacavir)

Protease Inhibitors (PI)

☐ IDV (Crixivan/indinavir)
 ☐ SQV-HSC (Invirase/saquinavir)
 ☐ SQV-SIS (Fortovase/saquinavir)

Therapy Options (10 of 63)

Therapy	Eff	Ad	Safety Considerations	File	Fills	Cost
<b>AZT-ddI-3TC-SQV-SGC</b>	1	1		q8h	26	\$43.46
<b>d4T, 3TC, NFV</b>				q8h	13	\$34.78
<b>AZT, 3TC, IDV</b>				q8h	10	\$32.24
<b>AZT, 3TC, NFV</b>				q8h	13	\$35.81
<b>d4T, 3TC, IDV</b>				q8h	10	\$31.20
<b>AZT, ddI, RTV, DL</b>				q8h	30	\$45.99
<b>ddI, d4T, IDV, NVP</b>				q8h	17	\$42.55
<b>d4T, 3TC, RTV</b>				q12h	16	\$38.46
<b>AZT, ddI, RTV, NV</b>				q12h	20	\$47.10

Show More

See All

Show Abstract for Retrovir

Show Abstract for Videx

Show Abstract for Epivir

Show Abstract for Fortovase

Show Therapy Study

Print Details for AZT-ddI-3TC-SQV-SGC

Print Top 10 Therapy Option Details

Print All Therapy Option Summaries

Print Top 10 Therapy Option Summaries

Hide Column "Eff"

Hide Column "Ad"

Hide Column "Safety Considerations"

Show Column "Met"

Show Column "Drug"

Hide Column "Firm"

Hide Column "Fills"

Hide Column "Cost"

Therapy Being Evaluated

None

WARNING::

CmtGenY, Con

Viral Load Testing required: Viral load testing should be repeated 21-35 days after initiation of, or a change of, antiRetroviral therapy to evaluate therapeutic efficacy and patient compliance. CmtGenY, Commentary65

Therapy Being Evaluated

None

WARNING::

CmtGenY, Con

Viral Load Testing required: Viral load testing should be repeated 21-35 days after initiation of, or a change of, antiRetroviral therapy to evaluate therapeutic efficacy and patient compliance. CmtGenY, Commentary65

Therapy Being Evaluated

None

WARNING::

CmtGenY, Con

Viral Load Testing required: Viral load testing should be repeated 21-35 days after initiation of, or a change of, antiRetroviral therapy to evaluate therapeutic efficacy and patient compliance. CmtGenY, Commentary65

Fig. 11E

- Therapy Initiation: Current treatment guidelines recommend initiation of antiRetroviral therapy for HIV-infected patients with HIV RNA (viral load) concentrations greater than 20,000 copies/ml (10,000 E<sub>q</sub>/ml bDNA) or CD4 counts less than 500 cells/ $\mu$ L (Ann. Int. Med., 1998). PrQualM, Commentary61
- Combination Therapy Recommended: Experts agree that the goal of antiRetroviral therapy should be to reduce the viral load to as low a level as possible for as long as possible. Initiation of therapy with a combination containing 2 nucleoside reverse transcriptase inhibitors (NRTI's) and a potent protease inhibitor have been shown to provide enhanced clinical benefit versus 2 drug combinations with regard to reduction in viral load and improved clinical outcomes. PrQualM, Commentary66

TPMS 11

How To

60

606 704

TPMS Patient

Medical History Chat Therapy Evaluation

CD4 (cells/cubic mm)

Viral Load (copies/mL)

1800

1700

1600

1500

1400

1300

1200

1100

1000

900

800

700

600

500

400

300

200

100

0

3TC

dAT

NVP

AZT

IDV

ddC

ddI

3TC

3TC

3TC

3TC

3TC

3TC

FIG. 12 A



A12

Phenotypic Resistance to 3TC from 3/15/1999 to present

12/1997 1/1998 2/1998 3/1998 4/1998 5/1998 6/1998 7/1998 8/1998 9/1998 10/1998 11/1998 12/1998 1/1999 2/1999 3/1999

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F18-12C

Medical History | Chart | Therapy Evaluation

<b>General</b>	
Patient ID: Features1	Birth Date: 1/1/1960
Physician: patient	Gender: Male
Weight (kg): 60.00	Date: 1/28/1999
Solid Dosage: Yes	Date: 1/28/1999
AIDS Diagnosis: <input type="checkbox"/> Date: <input type="checkbox"/>	
AIDS Defining Event: <input type="checkbox"/>	
Current ARV Therapy: <input type="checkbox"/> 3TC, AZT, NVP	
Non-ARV Drugs: <input type="checkbox"/>	
HIV Genotype: <input type="checkbox"/> L101P, M46I, V82A, V82AI, M41L, RT, Y181	
CD4 and Viral Load	
CD4 (cells/mm <sup>3</sup> ): 240	Specimen Date: 3/15/1999
Current Viral Load: 21500	VL Units: C/mL
Previous Viral Load: 2800	VL Units: C/mL
Hepatic Function	
AST/SGOT (IU/L): 25	Specimen Date: 1/28/1999
ALT/SGPT (IU/L): 25	Specimen Date: 1/28/1999
Renal Function	
BUN (mg/dL): 15.00	Specimen Date: 1/28/1999
Cr (mg/dL): 1.500	Specimen Date: 1/28/1999
Neutrophils	
Neutrophils (%): 15.00	Specimen Date: 1/28/1999
Neutropathy	
Neutropathy: No	Specimen Date: 1/28/1999
Pancreatitis	
Pancreatitis: No	Specimen Date: 1/28/1999
Hemoglobin	
Hemoglobin (g/dL): 15.00	Specimen Date: 1/28/1999
Hepatic Function	
AST/SGOT (IU/L): 25	Specimen Date: 1/28/1999
ALT/SGPT (IU/L): 25	Specimen Date: 1/28/1999
Renal Function	
BUN (mg/dL): 15.00	Specimen Date: 1/28/1999
Cr (mg/dL): 1.500	Specimen Date: 1/28/1999
Neutrophils	
Neutrophils (%): 15.00	Specimen Date: 1/28/1999
Neutropathy	
Neutropathy: No	Specimen Date: 1/28/1999
Pancreatitis	
Pancreatitis: No	Specimen Date: 1/28/1999
Hemoglobin	
Hemoglobin (g/dL): 15.00	Specimen Date: 1/28/1999

• NVPΔ Drug Interaction Alert: Patient is currently taking nifabutin and there is insufficient data to assess whether dose adjustments are necessary. These drugs should only be used in combination if clearly indicated and with careful monitoring. CmtDIP, Commentary33